

# I.B DESIGN OF THE TEM

## The TEM Top to Bottom:

- Electron gun
- Condenser lens(es)
- Lens aberrations
- Objective lens and specimen stage
- Projector lenses
- Camera and viewing system
- Vacuum system
- Electrical system

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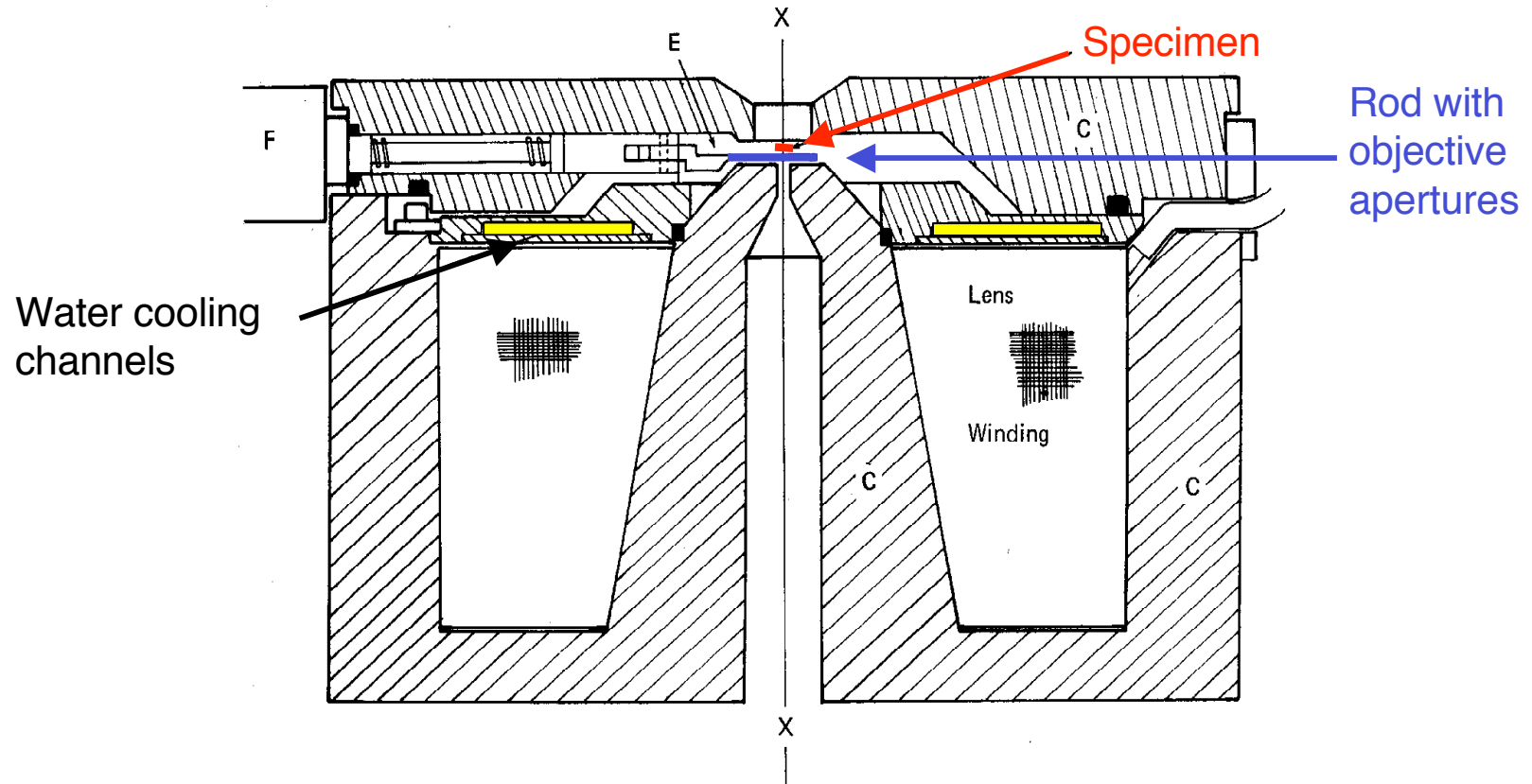
## I.B.4 Objective Lens and Specimen Stage

### **KEY CONCEPTS**

- Objective lens is **most critical** lens of TEM
- Performs the **first stage of imaging**
- Determines the instrument **resolving power** and **contrast**

# I.B.4 Objective Lens and Specimen Stage

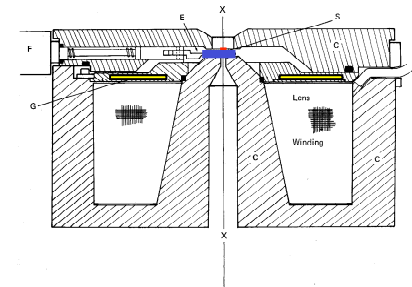
## I.B.4.b Objective Lens Construction



## I.B.4 Objective Lens and Specimen Stage

### I.B.4.b Objective Lens Construction

#### Requirements



- **Specimen** situated **close to** objective **front focal plane**
- **Focal length** should be as **small as practical** (minimizes chromatic & spherical aberration which decrease as focal length decreases)
- Specimen has to be **inside** the **lens field** to obtain the necessary short focal length (specimen in a confined space)
- Need **adequate clearance** for insertion of specimen, aperture, and anticontaminator
- Must provide for inserting devices (stigmators) to correct for asymmetries in the lens field

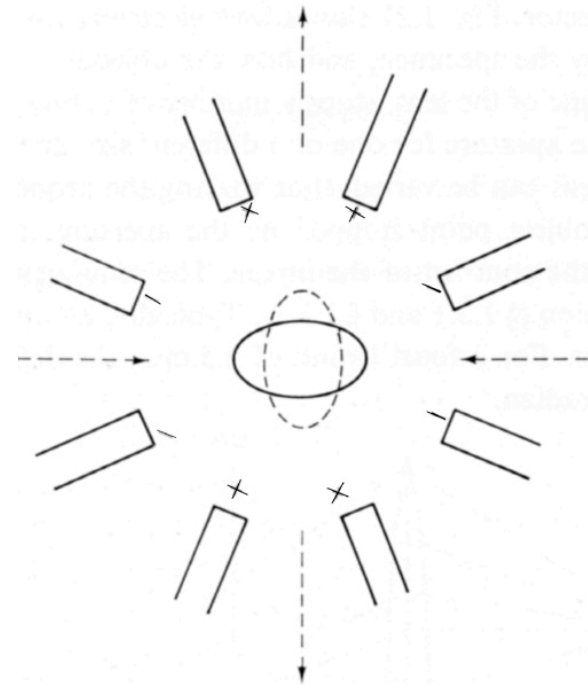
## I.B.4 Objective Lens and Specimen Stage

### I.B.4.c Objective Lens Asymmetry

- Impossible to manufacture lenses with perfectly rotationally symmetric magnetic fields about the optic axis
- If field is not perfectly symmetrical, the image will be **astigmatic** on as well as off the axis
- In older microscopes, **stigmators** consisted of iron pieces (**octupole**) that could be **physically moved** to make the field more symmetrical

**Octupole:** arrangement of magnets used to make the magnetic field of the objective lens symmetrical

- Modern microscopes use **electrostatic fields** to correct lens astigmatism



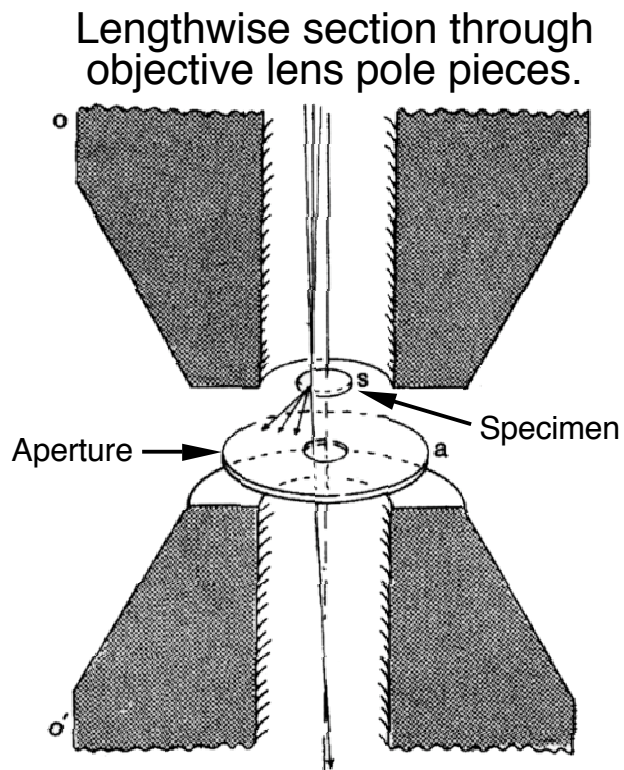
From Agar, Fig. 1.22, p.28

## I.B.4 Objective Lens and Specimen Stage

### I.B.4.e Objective Aperture

**FUNCTION:** Intercepts  $e^-$  which have been **scattered** by the specimen through **large angles**

**POSITION (most common):** **Back focal plane** of the objective lens



- Does not restrict field of view
- Contamination effects reduced in this position since only widely scattered  $e^-$  strike the periphery of the aperture opening

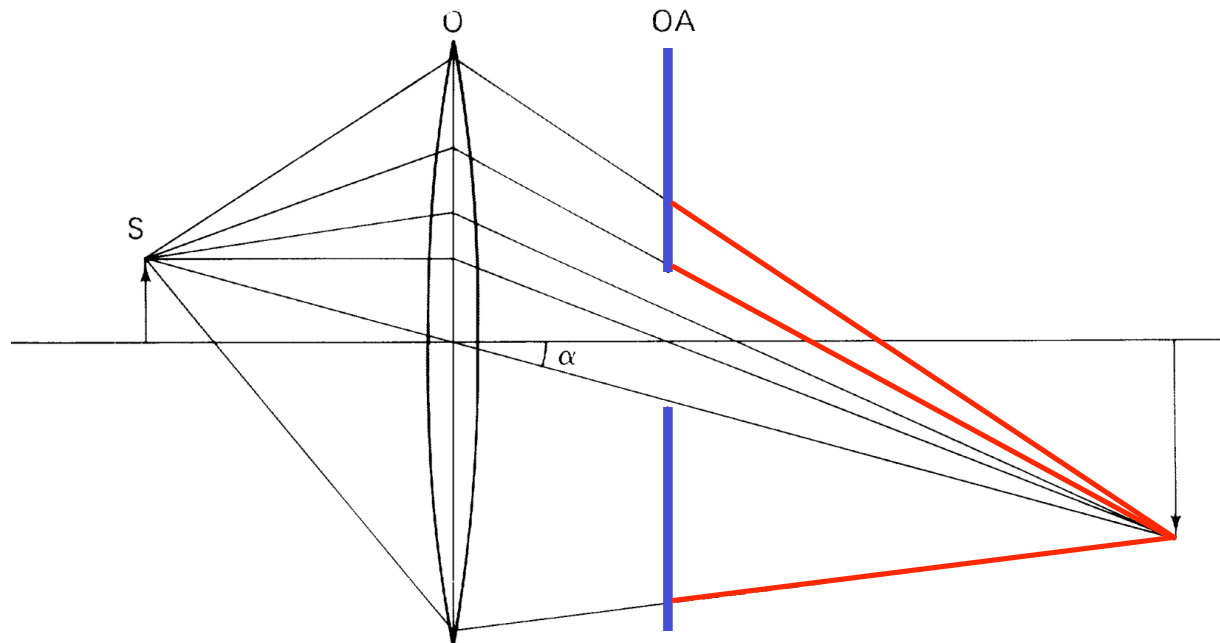
## I.B.4 Objective Lens and Specimen Stage

### I.B.4.e Objective Aperture

**FUNCTION:** Intercepts  $e^-$  which have been **scattered** by the specimen through **large angles**

**POSITION (most common):** **Back focal plane** of the objective lens

In this position, the objective aperture screens out widely scattered electrons from being imaged





## I.B.4 Objective Lens and Specimen Stage

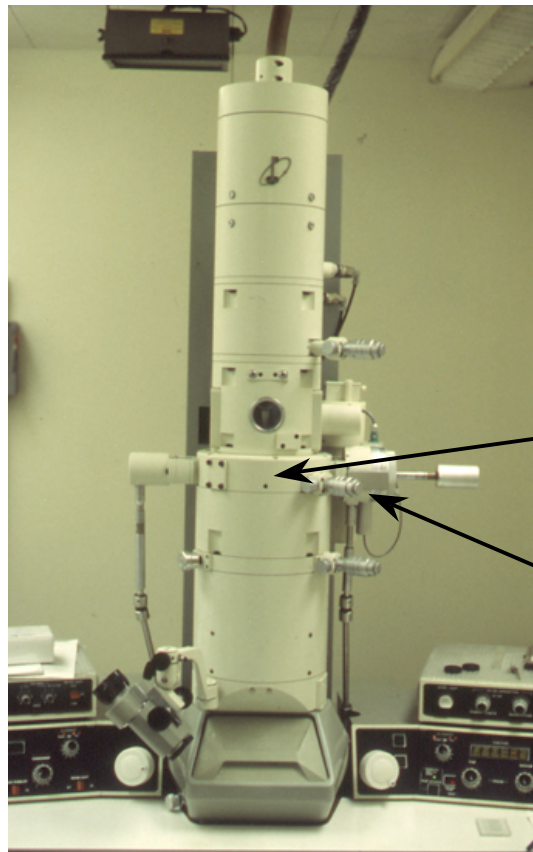
### I.B.4.e Objective Aperture

#### *Factoids, Factoids, and even more Factoids:*

- ~ **25-75 $\mu\text{m}$**  hole diameter
- **Smallest** apertures give **best image contrast**
- Must be **perfectly circular** and **clean** (or imaging field will be distorted)  
Contaminated aperture can act like a weak electrostatic lens causing image astigmatism
- Difficult to manufacture with good circular symmetry
- Contamination effects more serious with smaller apertures
- Use **ultrathin, self-cleaning** metal apertures
- Older TEMs: platinum or molybdenum apertures (**need regular cleaning**)

## I.B.4 Objective Lens and Specimen Stage

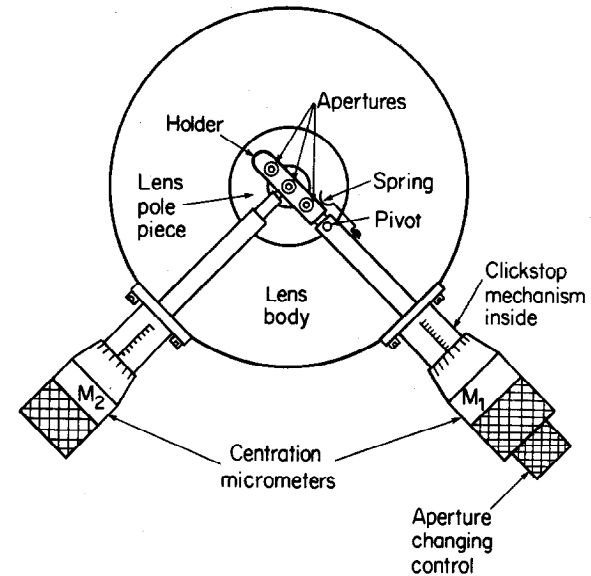
### I.B.4.e Objective Aperture



Objective lens

Objective aperture

Most TEMs have an aperture holder with space for 3 apertures



From Meek, 1st ed., Fig. 5.2, p.95

### Aperture centering (simplest method):

Image the **back focal plane** of the objective lens on the fluorescent screen (TEM in SA diffraction mode)

## I.B.4 Objective Lens and Specimen Stage

### I.B.4.f Specimen Stage

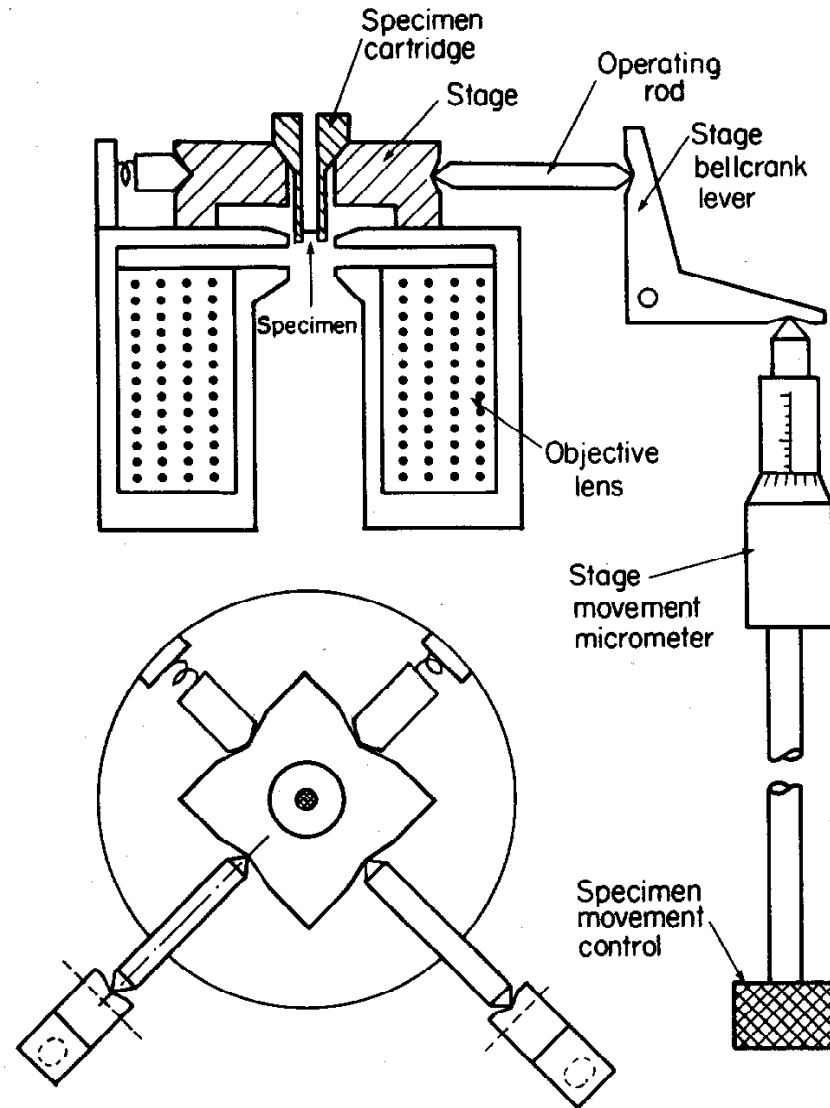
#### Requirements of Suitable Stage

- Allow simple and rapid specimen exchange
- Must have a specimen **airlock**
- Should sit in a plane well defined with respect to its position along the axis of the optical system
- Provide **minimum backlash** (<100 nm) - **no drift**
- Provide minimal vibrations, thermal motions, mechanical drift, and movements
- Make **good thermal contact** with specimen

# I.B.4 Objective Lens and Specimen Stage

## I.B.4.f Specimen Stage

### Top Entry Stage



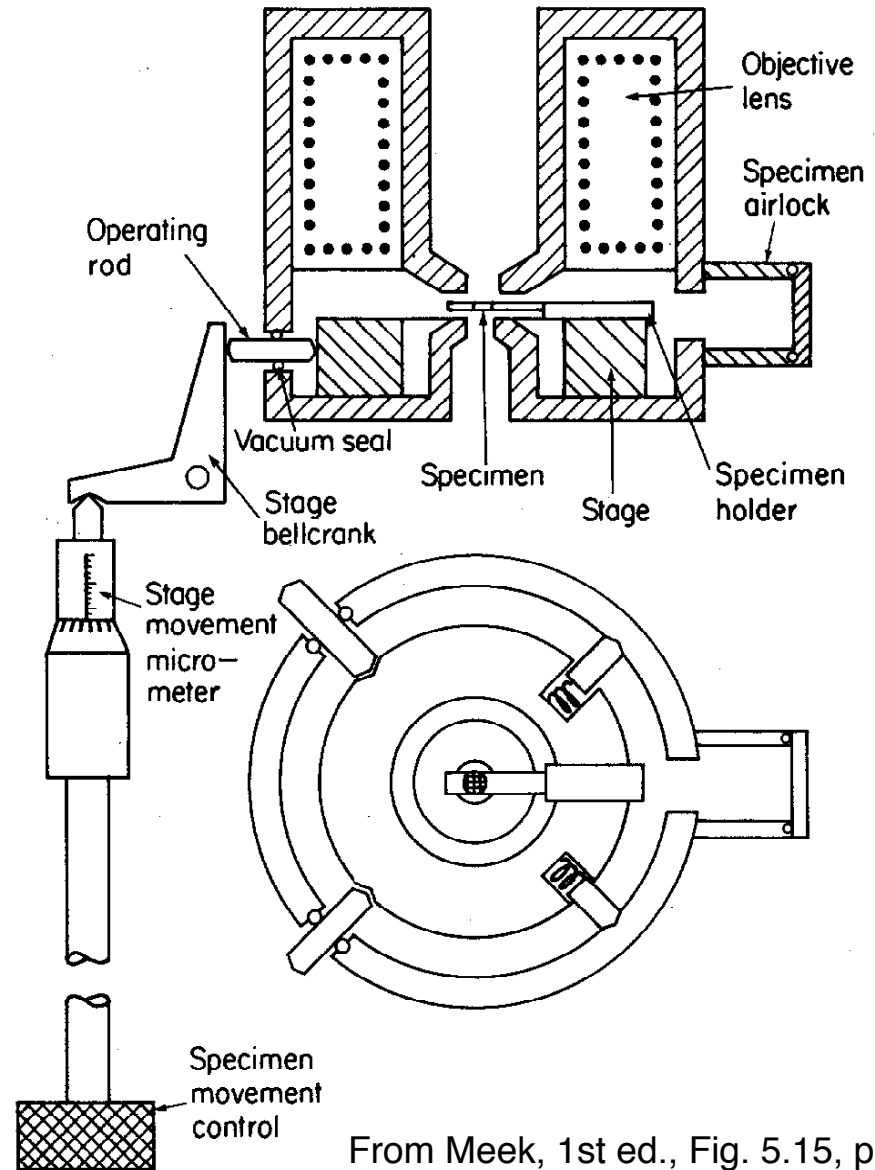
Drop-in cartridge type of objective lens, stage, and stage motion.

# I.B.4 Objective Lens and Specimen Stage

## I.B.4.f Specimen Stage

### Side Entry Stage

The side-specimen-entry, immersion type of objective lens, stage, and stage motion



From Meek, 1st ed., Fig. 5.15, p.115

## I.B.4 Objective Lens and Specimen Stage

### I.B.4.g Special Stages

**One for nearly every need!!!**

- Tilt stage
- Multiple specimen stage
- Furnace heating stage
- Grid heater stage
- Cold stage
- Straining stage
- Gas reaction stage
- Hydration or 'wet' stage
- Many, many more....

See lecture notes (p.48) for more details

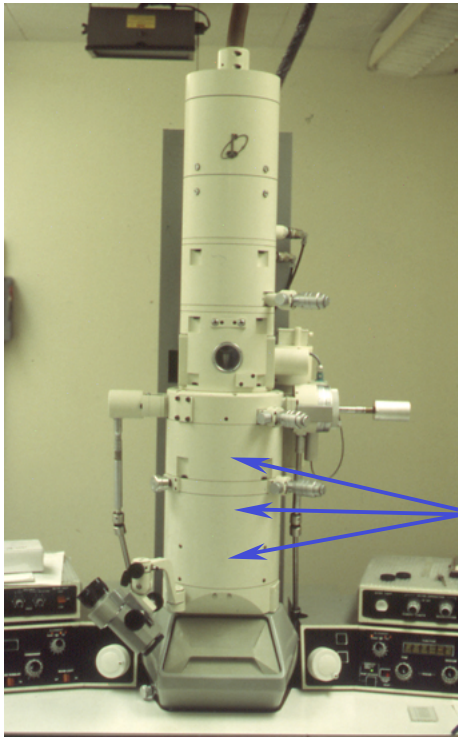
End of Sec.I.B.4

## I.B.5 Projector Lenses

### I.B.5.a Description

## Projector Lens Systems

Produce and control magnification of the final image



Projector lenses

- Modern instruments use 3 or 4 projector lenses (diffraction, intermediate, and one or two projector lenses) to give a wide magnification range
- Each microscope has a **different formula** for producing a wide range of image magnifications (typically  $\sim 1000X$  up to  $>500,000X$ )
- **Object** of 1st projector lens: **intermediate image produced by objective lens**
- Last projector lens magnifies an area of the 2nd or 3rd intermediate image that may be several millimeters in diameter



## I.B.5 Projector Lenses

### I.B.5.b Distortion

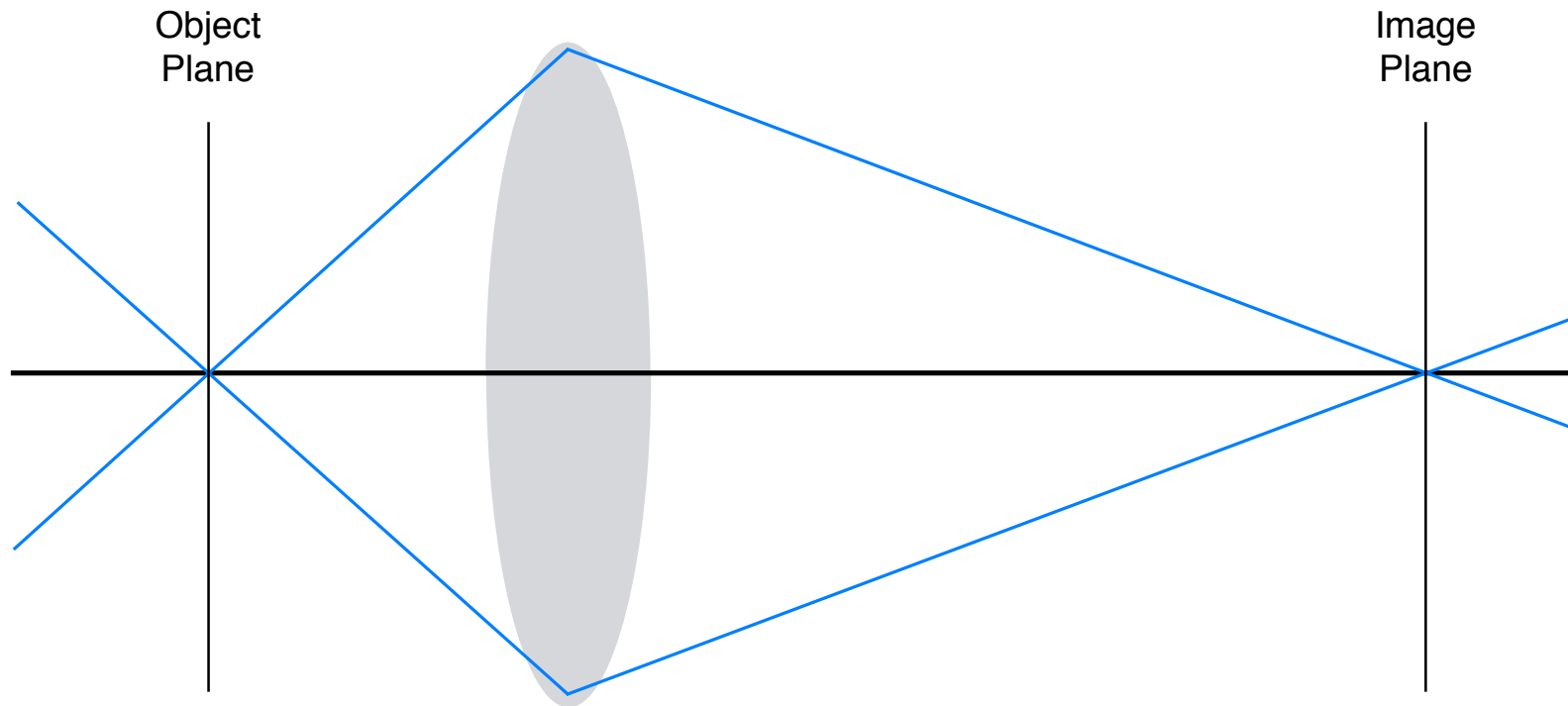
- Aberrations in the projector lenses **do not influence resolution of final image** but may produce some image distortion (barrel, pincushion, spiral)

See lecture notes (pp.37-39 and p.49) for details

# I.B.5 Projector Lenses

## I.B.5.c Depth of Field and Depth of Focus

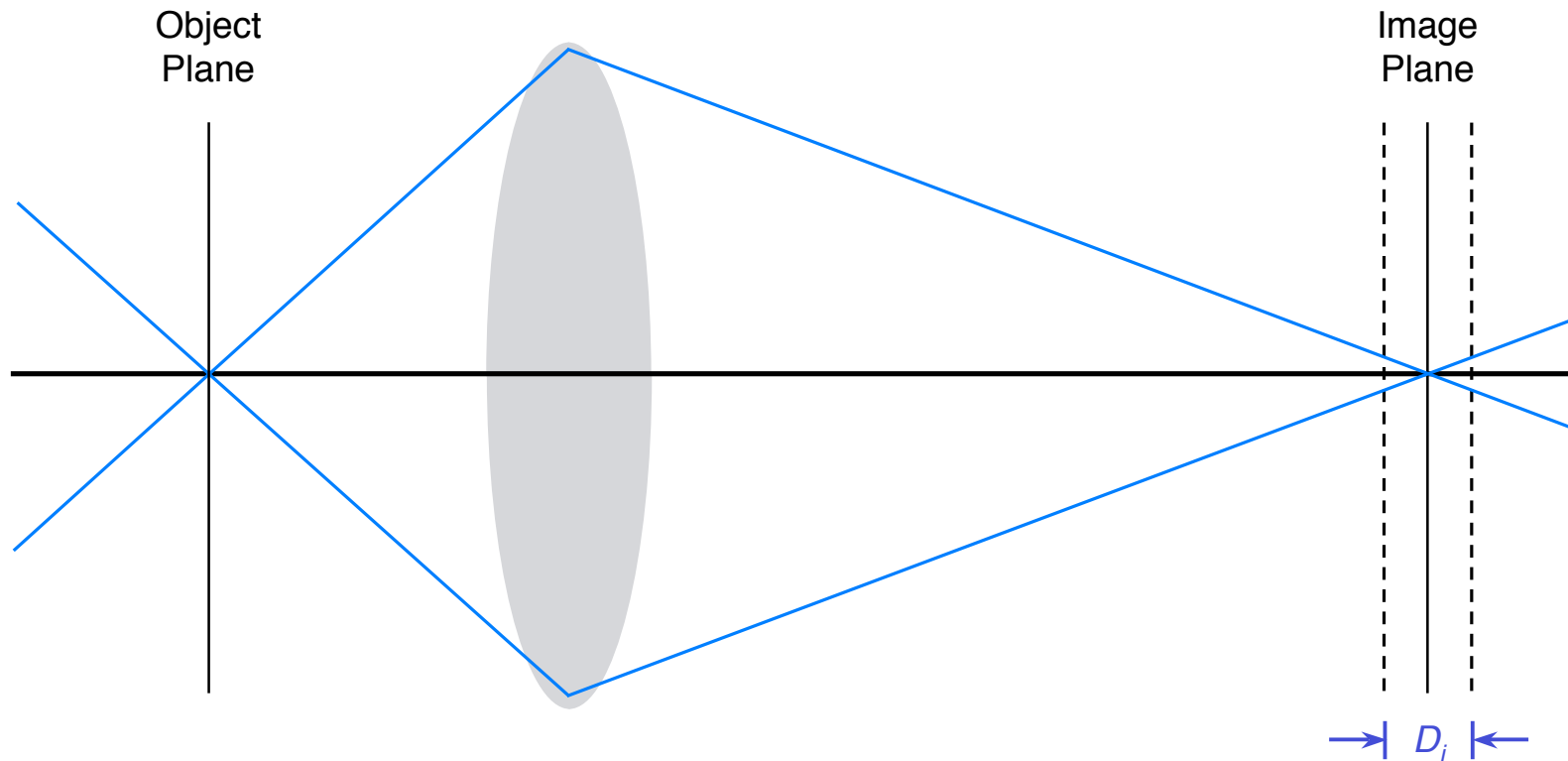
**Recall:** Any lens, however perfect, can only image a point object as an Airy disc, the diameter of which = lens resolving power



## I.B.5 Projector Lenses

### I.B.5.c Depth of Field and Depth of Focus

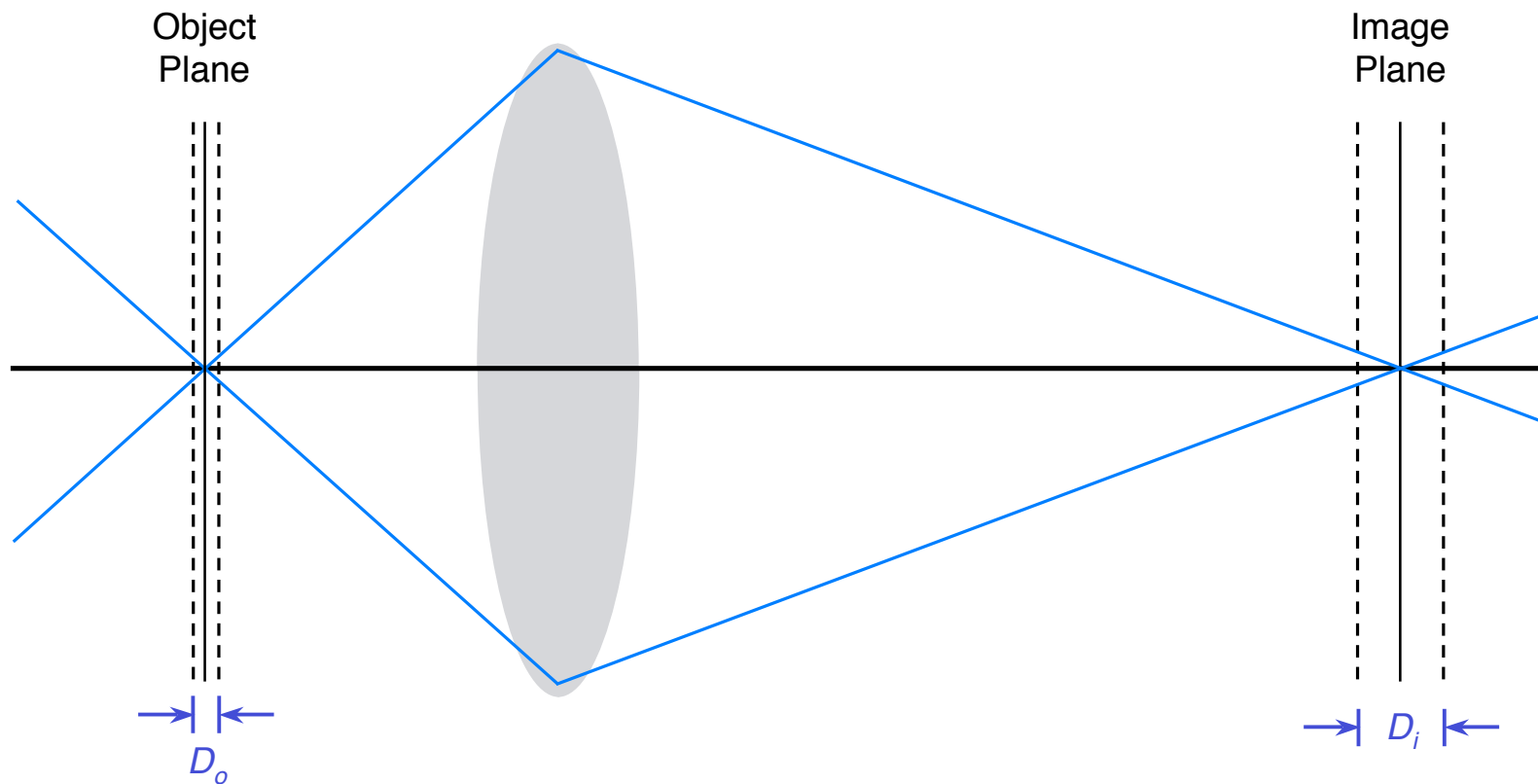
There is a finite distance along the axis,  $D_i$ , where the image appears equally sharp. This distance is called **depth of focus**.



## I.B.5 Projector Lenses

### I.B.5.c Depth of Field and Depth of Focus

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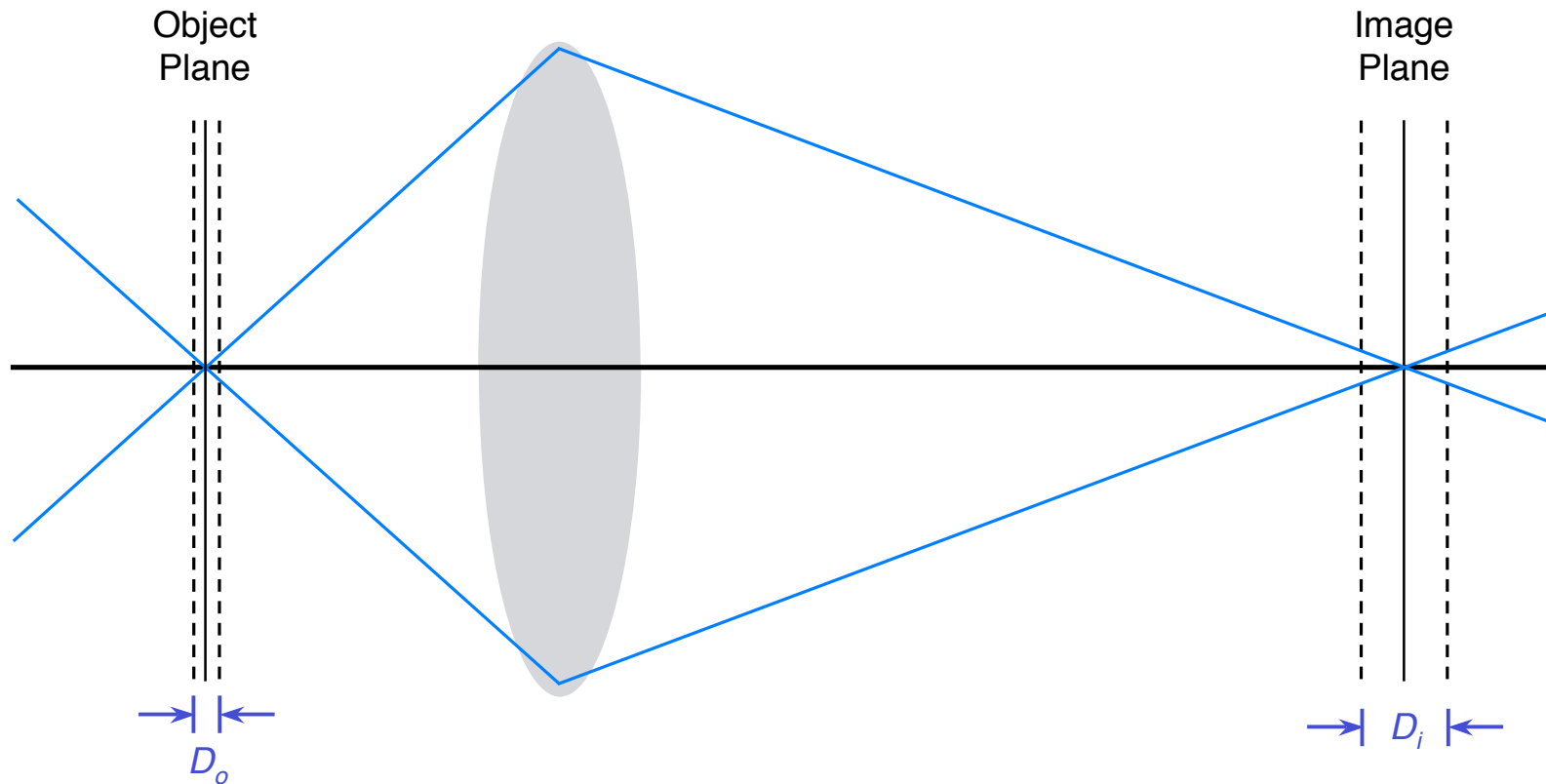


An analogous distance,  $D_o$ , along the axis on the **object side** over which the object could be moved and still give a maximally sharp image (**at position of "exact" image plane**). This distance is called the **depth of field**.

## I.B.5 Projector Lenses

### I.B.5.c Depth of Field and Depth of Focus

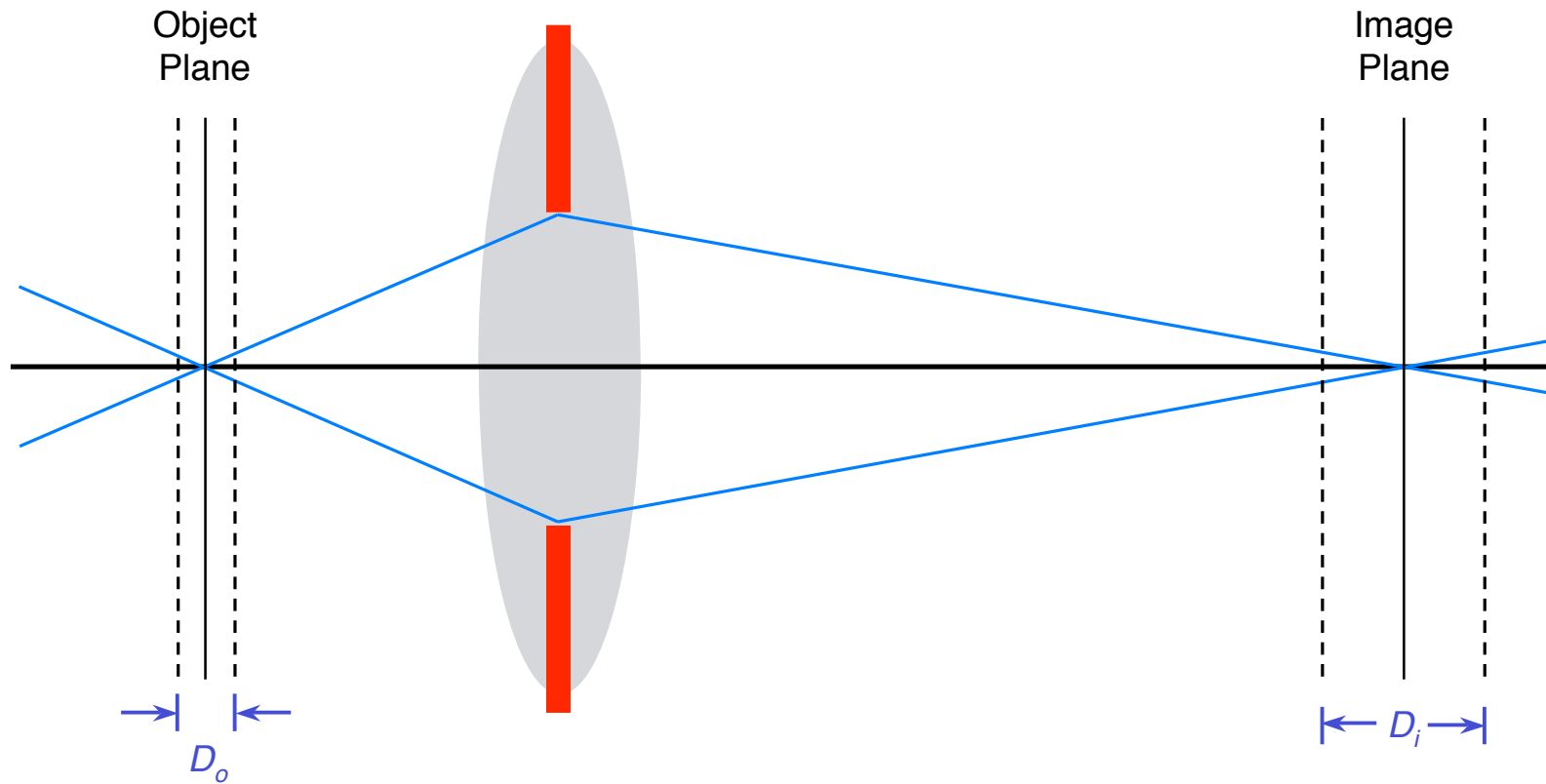
*Decreasing* the aperture of the lens *increases both*  $D_o$  and  $D_i$ .



## I.B.5 Projector Lenses

### I.B.5.c Depth of Field and Depth of Focus

*Decreasing* the aperture of the lens *increases both*  $D_o$  and  $D_i$ .



## I.B.5 Projector Lenses

### I.B.5.c Depth of Field and Depth of Focus

#### Depth of Field (Object/Specimen Plane)

$$D_o = \frac{2d}{\tan \alpha_o}$$

$d$  = minimum object spacing one hopes to resolve

$\alpha_o$  = **objective lens** semi-angular aperture

**EXAMPLE:** For  $d = 1.0$  nm and  $\alpha_o = 5 \times 10^{-3}$  radians,  $D_o = 400$  nm  
(thicker than most TEM specimens)

**CONSEQUENCE:** A **thin specimen** appears **equally sharp** through its  
**entire thickness**

## I.B.5 Projector Lenses

### I.B.5.c Depth of Field and Depth of Focus

#### Depth of Focus (Image Plane)

$$D_i = \frac{M^2 2d}{\tan \alpha_o} = D_o M^2$$

$M$  = **total** magnification of compound magnifying system

$d$  = minimum object spacing one hopes to resolve

$\alpha_o$  = **objective** lens semi-angular aperture

**EXAMPLE:** If  $M = 50,000X$ ,  $d = 1.0 \text{ nm}$ , and  $\alpha_o = 5 \times 10^{-3}$  radians, then  
 **$D_i = 1000 \text{ meters!!!}$**



## I.B.5 Projector Lenses

### I.B.5.c Depth of Field and Depth of Focus

#### **Consequences (Key Concepts):**

- Fluorescent screen, photographic plate or film can be placed **ANYWHERE** on the optic axis **beneath** the projector lens and the final image **will be in equally sharp focus** (though **magnification WILL** differ)
- Large depth of field/focus does **NOT** eliminate the requirement for **VERY CAREFUL FOCUSING** of the image (by adjusting the objective lens strength)

Ugh.....just when you thought something might be easy!

## I.B.5 Projector Lenses

### I.B.5.c Depth of Field and Depth of Focus

#### **Factoids:**

- **LM:** depth of field and depth of focus are roughly the same magnitude
- **TEM:** depth of **focus** is **many times greater** than the depth of field (by the factor =  $M^2$ )

End of Sec.I.B.5

# I.B DESIGN OF THE TEM

## I.B.6 Camera and Viewing System

### I.B.6.a Viewing the Image

- Electron optical image is projected onto a **fluorescent screen**
- **Fluorescent screen**: surface coated with a layer of activated zinc sulfide crystals
- **Kinetic energy of electrons** in the image is **transformed into light energy** through fluorescence
- **Resolution of image on screen** determined by size of these crystals ( $\sim 50\text{-}75\ \mu\text{m}$ )

## I.B.6 Camera and Viewing System

### I.B.6.b Photographing the Image

- Photographic recording done at **magnification sufficient to maintain resolution** in electron image
- **Resolution of photographic emulsions** generally ~ 4-5 times better than the fluorescent screen
- **Exposure time** may be determined from a reading taken by a photo-cell looking at the fluorescent screen or by reading the current on the screen itself
- Most common types of photographic materials are 3x4" sheets of film, 35mm film, and 70mm roll film

## I.B.6 Camera and Viewing System

### I.B.6.c Photographic Emulsion

- **Photographic recording material:** generally a plastic base coated with an emulsion
- **Emulsion** = layer of gelatin containing photosensitive silver halide
- The fine-grained negative (electron micrograph) contains a more detailed and higher contrast image than that seen on the fluorescent screen

**NOTE:** J. Turek will give more details about photography of electron images. See also lecture notes, pp.86-93.

End of Sec.I.B.6

# I.B DESIGN OF THE TEM

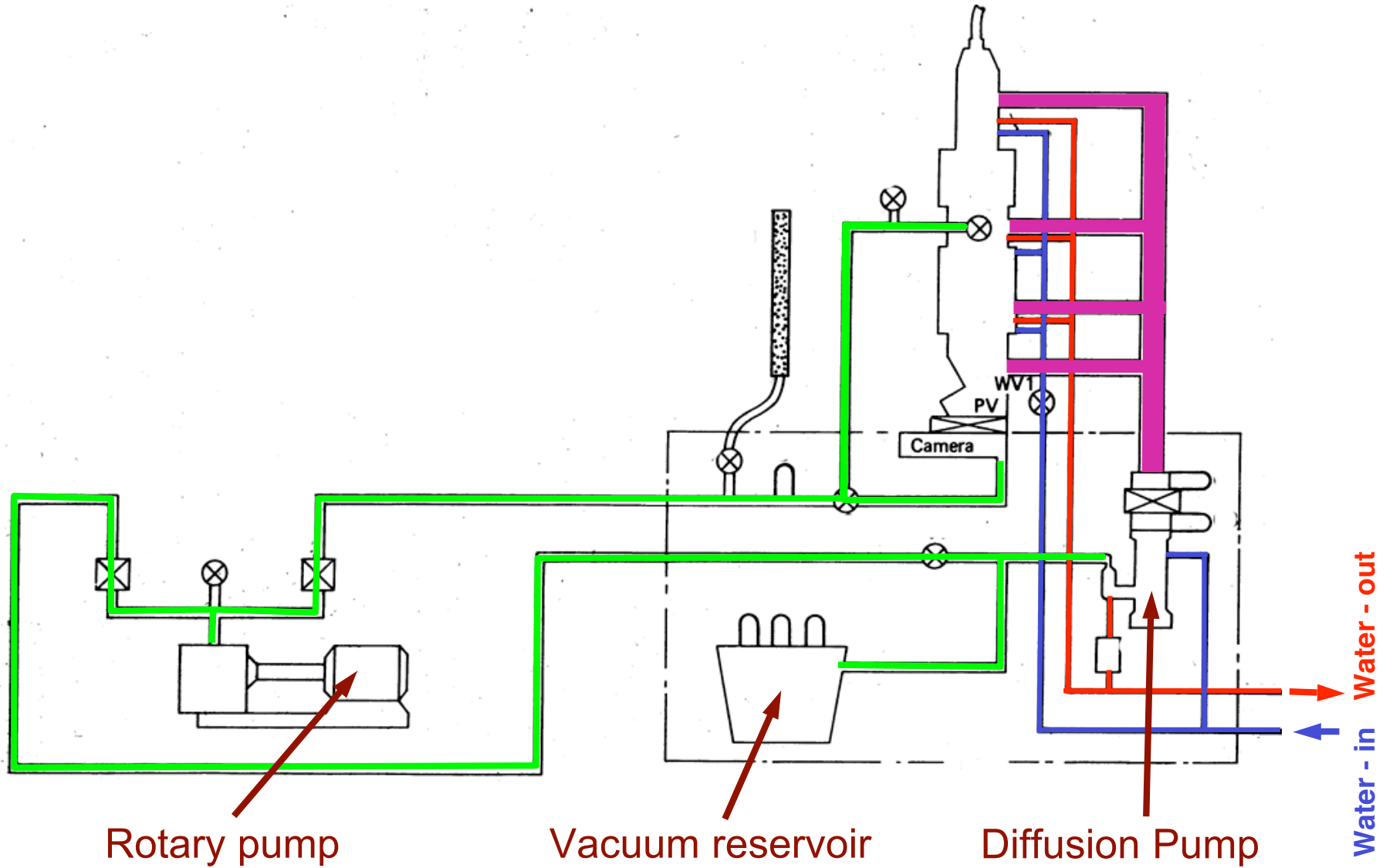
## I.B.7 TEM Vacuum Systems

### KEY CONCEPTS

- High vacuum needed in order to allow passage of e<sup>-</sup> through microscope
- **Mean free path** of an electron at atmospheric pressure (760 Torr) is a measly  $6.5 \times 10^{-6}$  cm (**65 nm!**)
- TEMs operate at  **$10^{-6}$  Torr** or lower (e- mean free path is **> 50 meters!**)
- Most TEMs have at least **two types of pumps in tandem**



# I.B.7 TEM Vacuum Systems



## I.B.7 TEM Vacuum Systems

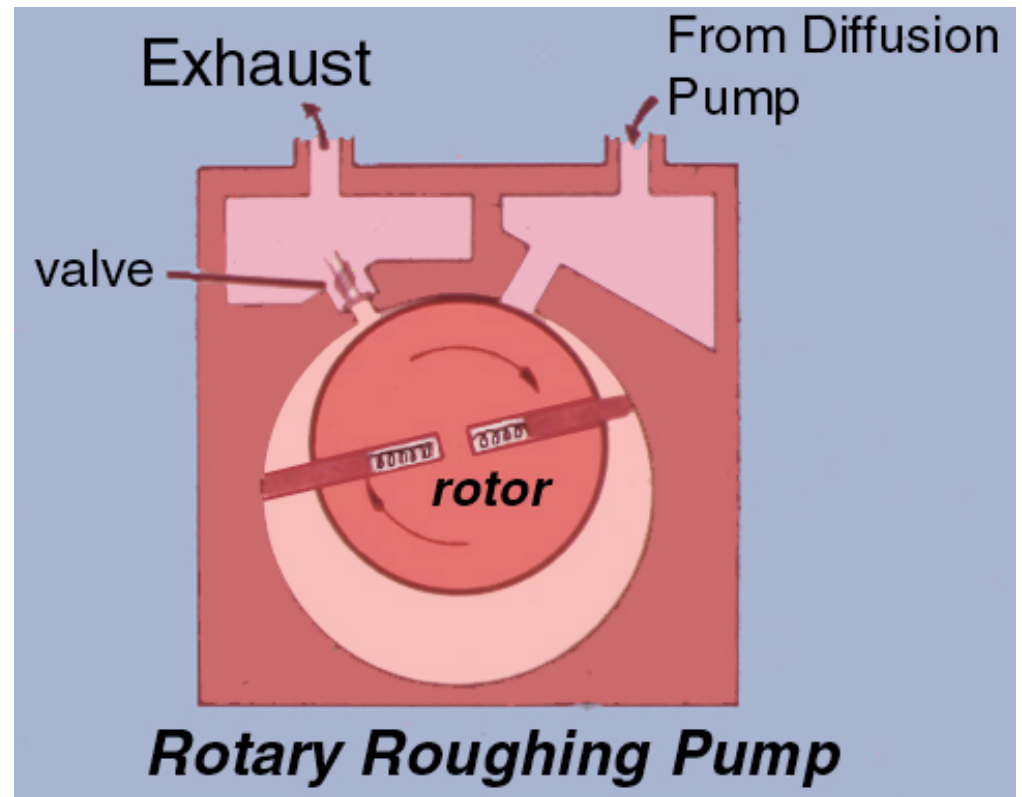
### I.B.7.b Types of Pumps

## Rotary Vane Pump

Mechanical pump that compresses and displaces trapped gases through the action of sliding vanes in a non-symmetrical chamber.

Pull a vacuum of  $10^{-3}$  to  $10^{-4}$  Torr

Often referred to as a **roughing pumps** since they do the initial “rough” pumping down from atmospheric pressure.



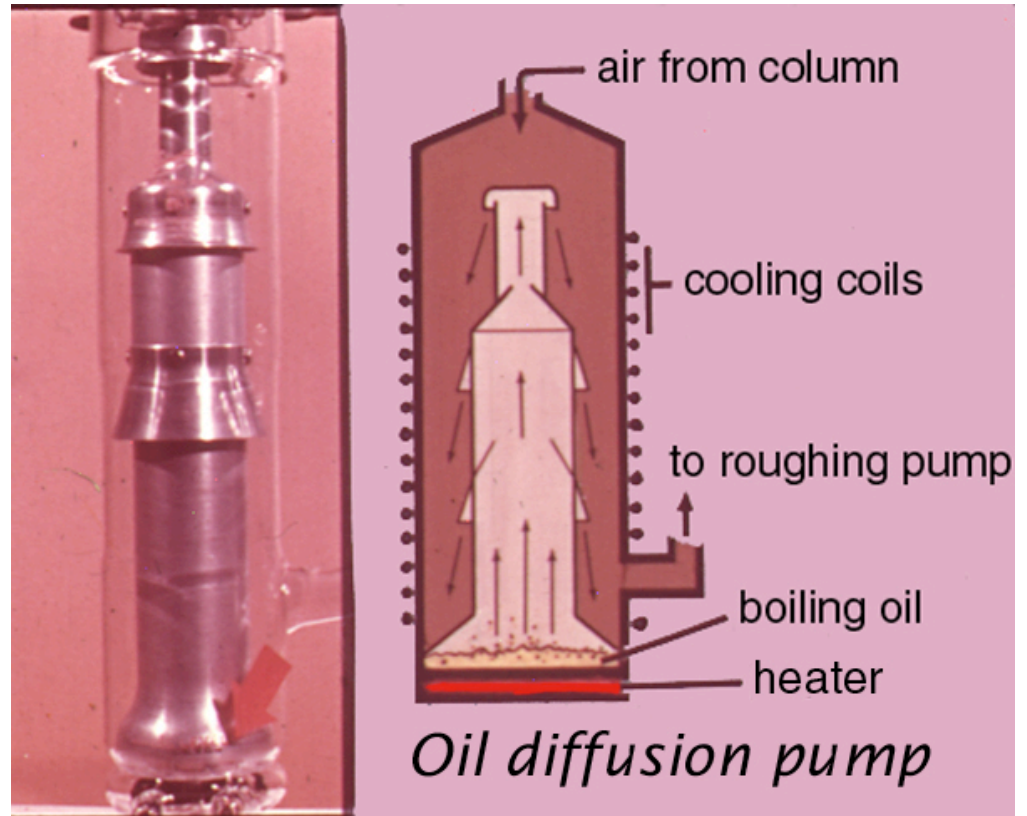
From Crang and Ward

## I.B.7 TEM Vacuum Systems

### I.B.7.b Types of Pumps

# Diffusion Pump

- Use **hot oil vapor** moving at supersonic speed to trap air molecules and remove them from the microscope.
- Diffusion pumps **need to be backed up by rotary pumps** since they require a backing pressure of at least  $2 \times 10^{-1}$  Torr to operate. They also require a water cooling source.
- Develop a vacuum of  **$10^{-5}$  to  $10^{-7}$  Torr**. **High vacuum** is defined as  $10^{-3}$  to  $10^{-7}$  Torr. **Ultra-high vacuum** is below  $10^{-7}$  Torr.



From Crang and Ward

## I.B.7 TEM Vacuum Systems

### I.B.7.b Types of Pumps

#### **Ion Pump**

- Vacuums as high as  $10^{-9}$  Torr can be obtained by ionizing the particles to be out gassed in a strong electric field and subsequently causing them to be trapped at a surface.
- Many modern TEMs (*e.g.* FEI CM200 and CM300 FEG) use **ion getter pumps** to reduce the vacuum in the gun and specimen area to less than  $10^{-7}$  Torr, thereby increasing filament lifetime and reducing specimen contamination

## I.B.7 TEM Vacuum Systems

### I.B.7.b Types of Pumps

#### **Cryo Pump**

If a **surface** is cooled below the condensation temperature of a vapor, it acts as an **effective pump**, since vapor reaching the surface cannot escape again.

**Liquid nitrogen cooled surface** is very effective for trapping organic vapors, but not lighter gases.

#### **Other Pump Types**

**Turbomolecular Pump:** An axial flow turbine pump consisting of alternating circular rotor and stator disks with inclined blades.

These pumps do not use oil vapor to establish high vacuum and **may be used alone**.

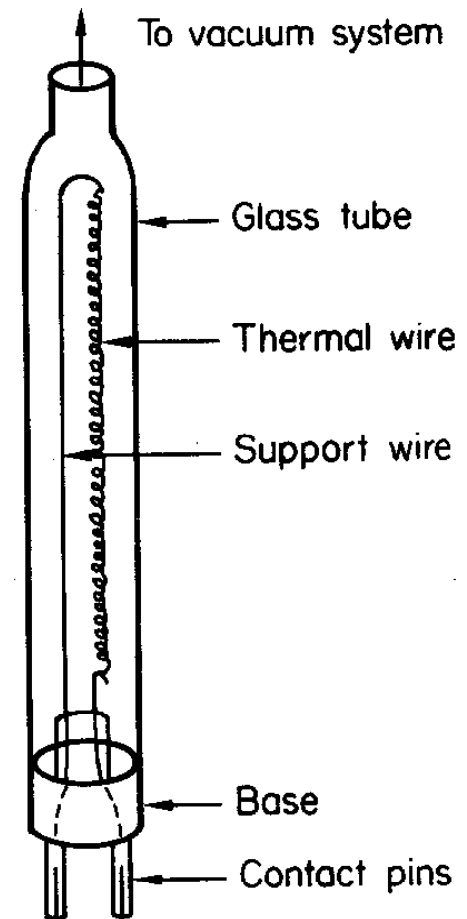
## I.B.7 TEM Vacuum Systems

### I.B.7.b Measuring Vacuum (Gauges)

Need to **monitor the quality** of the vacuum obtained by each type of pump

**Pirani Gauge:** Uses thermal conductivity to measure vacuum

- Heat dissipation from a heated filament is function of gas pressure
- Operate effectively from atmospheric pressure to  $\sim 10^{-4}$  Torr



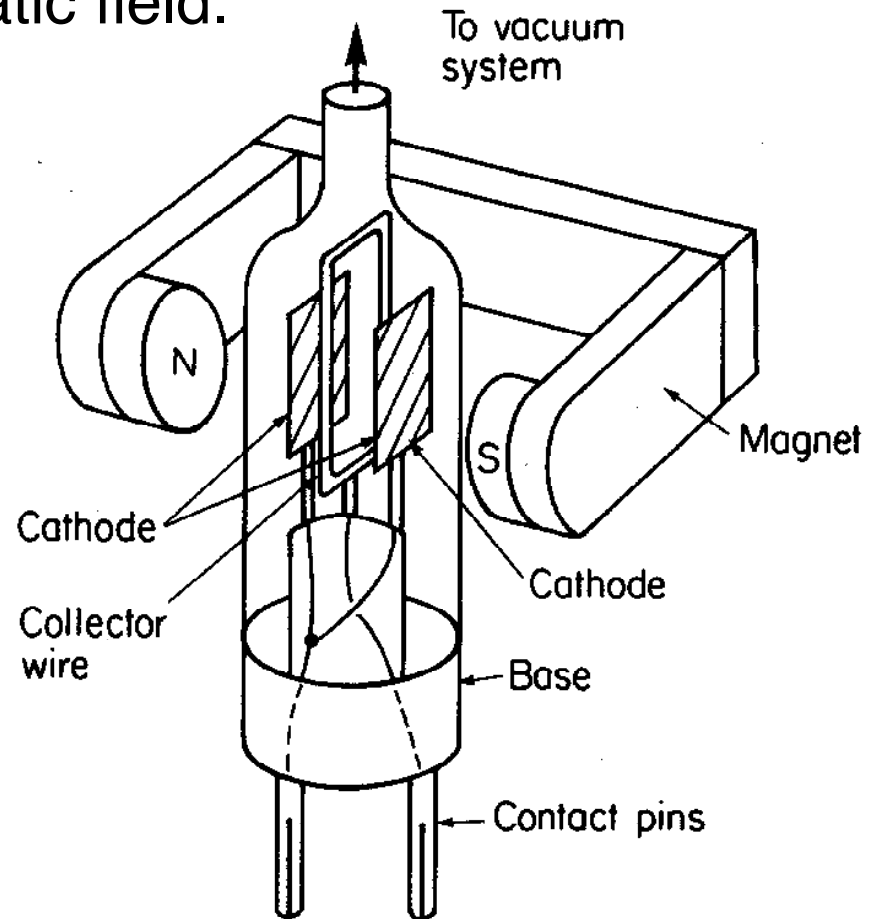
From Meek, 1st ed., Fig.6.3a, p.140

## I.B.7 TEM Vacuum Systems

### I.B.7.b Measuring Vacuum (Gauges)

**Penning (Ion) Gauge:** Depends on ionization of the gas molecules in a high electrostatic field.

- Electric field is developed between 2 cathodes and an anode which is placed between them.
- Effective from  $\sim 10^{-4}$  Torr and at higher vacuums



From Meek, 1st ed., Fig.6.4a, p.142

## I.B.7 TEM Vacuum Systems

### Limits to High Vacuum

- Pumping system will eventually come to an **equilibrium state** that is less than what is possible theoretically
- Air molecules stick to metal surfaces and are constantly being released
- Lint and dirt on O-rings leads to air leaks
- Photographic materials constantly leak water vapor, plasticizers, and other volatiles
- **Grease on gaskets and O-rings** are a source of hydrocarbon contamination
- **Grease from fingerprints** (*e.g.* on the specimen holder) contaminates everything in the vacuum
- **Cardinal Rule: Wear gloves** whenever handling any part exposed to the vacuum



## I.B.7 TEM Vacuum Systems

### Problems Caused by Poor Vacuum

**Contamination of specimen:** Hydrocarbon residue from oil, grease etc. gets hit by the  $e^-$  beam and decomposes to H and C. C atoms get fixed to the specimen and destroys resolution.

**Decreased filament life** due to etching of the filament

**Beam can strip or etch** the specimen. Residual water in the specimen is ionized and  $\text{OH}^-$  ions attack carbon in the specimen to make  $\text{CO}$ . This essentially 'burns' up the specimen.

Some of these problems are reduced or eliminated by use of an **anticontaminator**, which is a liquid  $\text{N}_2$  cooled metal ring positioned near the specimen that traps contaminants before they reach the specimen.

End of Sec.I.B.7

# I.B DESIGN OF THE TEM

## I.B.8 Electrical System

### I.B.8.a General Requirements

#### **TEM electrical power supplies must provide:**

- **Current to heat** the **filament** and generate image-forming electrons
- **High voltage** to accelerate the emitted e<sup>-</sup> beam.
- **Current to each magnetic lens** to provide the necessary focusing magnetic fields.
- **Power to many circuits** such as stigmators, beam deflectors, camera, and camera shutter, exposure meter, focus wobbler, safety devices, relay switches, heaters of the diffusion pumps, the rotary pump, etc.

## I.B.8 Electrical System

### I.B.8.b Filament Current Supply

- Filament requires between **2.5-3 amps** as a heater current
- Supply is usually D.C. to avoid ripples that would modulate the e<sup>-</sup> beam

### I.B.8.c High Voltage Supply

- Must supply ~ 0.1 milliamperes or less
- Accelerating potential supply is **high voltage and low current** (delivers 2-10 watts of power)

## I.B.8 Electrical System

### I.B.8.d Lens Current Supply

- Individual current supplies are required for each lens with **stabilities** of a **few parts per million**
- Supplies transistorized and operate at relatively low voltages. When high voltage to the gun is changed, a corresponding change is made in the lens currents to **maintain focus and magnification**
- Control of lens currents is programmed into the TEM so the correct lens combinations are obtained for minimizing distortion

## I.B.8 Electrical System

### I.B.8.e Stability Requirements

- For **high voltage stabilization**, stable high gain amplifiers with negative feedback from a resistive voltage divider between the cathode and ground are used
- High voltage generator is **enclosed in an oil-filled tank** (air not a good insulator)
- **Voltage** needs to be stable within **few parts per million**

## I.B.8 Electrical System

### I.B.8.e Stability Requirements

#### **Lens Current Stabilization**

- **Objective lens** has the **strictest stability requirement** of all the lenses (one part in  $10^5$ )
- Stigmators and beam deflectors require stabilities of same magnitude as lenses and are generally fed from the lens circuits
- If current through any imaging lens varies, the image rotates about a point called the **current rotation center** (coincident with the viewing screen center if the microscope is properly aligned)
  - Micrograph taken under these circumstances will be blurred at the edges and sharp at the center
- Fluctuating accelerating voltage gives rise to changes in magnification
  - Image grows or contracts radially about a point called the **voltage center** which should be coincident with the center of the viewing screen

## I.B.8 Electrical System

### I.B.8.e Stability Requirements

## Lens Current Stabilization

**Recall:** focal length of a magnetic lens is a function of the square of the current passing through the lens

$$f = \frac{KV_r}{(N \cdot I)^2}$$

- Hence, **current stability requirement** for an electromagnetic lens is higher than that of the accelerating voltage (by about a factor of 2)
- If, for example, accelerating voltage must be stable to one part in  $5 \times 10^4$ , then the lens current must be stabilized to within one part in  $10^5$  over the period required for recording the image (1-5 seconds)



End of Sec.I.B.8